What is claimed is:

1. A process for the preparation of a compound of formula V

$$R^1 \longrightarrow O \longrightarrow O - SO_2R^2 \qquad V$$

wherein R¹ is aryl or heteroaryl, and R² is lower alkyl, aryl or trifluoromethyl;

comprising brominating a compound of formula VI,

$$OOO$$
 OO OO OO OO

wherein R³ is lower alkyl, condensing the resulting brominated compound with R¹C(O)NH₂, wherein R¹ is as above, to form a compound of formula VII,

$$R^1 \underbrace{\hspace{1cm}}^{O} \underbrace{\hspace{1cm}}_{CO_2R^3} VII$$

wherein R¹ and R³ are as above,

reducing the compound of formula VII to convert the ester group to a corresponding alcohol, and

introducing a $-SO_2R^2$ group, wherein R^2 is as above, onto the reduced compound of formula VII to yield the compound of formula V.

2. A process for the preparation of a compound of formula V,

$$R^1 \longrightarrow 0$$
 $O \longrightarrow SO_2R^2$
 V

wherein R¹ is aryl or heteroaryl, and R² is lower alkyl, aryl or trifluoromethyl;

comprising brominating a compound of formula VI

wherein R³ is lower alkyl,

converting the brominated compound to a compound of formula X,

wherein R^3 is as above and R^4 is lower alkyl, lower-alkyl-carbonyl, lower-alkoxy-carbonyl, aryl-carbonyl, $P(O)(OR^5)_2$, or $Si(R^6)_3$, wherein each R^5 independently represents lower alkyl or aryl, and each R^6 independently represents lower alkyl or aryl;

subsequently condensing the compound of formula X with an amide $R^1C(O)NH_2$, wherein R^1 is as above, to obtain a compound of formula VII,

$$R^{1}$$
 $CO_{2}R^{3}$ VII

wherein R¹ and R³ are as above,

reducing the compound of formula VII to convert the ester group to a corresponding alcohol and

subsequently introducing a $-SO_2R^2$ group, wherein R^2 is as above, to yield said compound of formula V.

- 3. A process according to claim 2, wherein R³ is methyl or ethyl.
- 4. A process according to claim 2, wherein R² is methyl, ethyl, trifluoromethyl or 4-methylphenyl.
- 5. A process according to claim 4, wherein R² is methyl.
- 6. A process according to claim 2, wherein R^1 is phenyl.
- 7. A process according to claim 2, wherein R¹ is thiophen-2-yl.
- 8. A process for the preparation of 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedione or Sodium 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedionate comprising the steps:
 - a) reacting methyl- or ethyl 3-oxovalerate with bromine to yield methyl- or ethyl 4-bromo-3-oxovalerate,
 - b) reacting the methyl- or ethyl 4-bromo-3-oxovalerate with benzamide to yield methyl- or ethyl 2-(5-methyl-2-phenyl-4-oxazolyl)acetate,
 - c) converting the methyl- or ethyl 2-(5-methyl-2-phenyl-4-oxazolyl)acetate to 2-(5-methyl-2-phenyl-4-oxazolyl)ethanol,

- d) reacting the 2-(5-methyl-2-phenyl-4-oxazolyl)ethanol with methanesulfonylchloride to yield 2-(5-methyl-2-phenyl-4-oxazolyl)ethanol methansulfonyl ester,
- e) reacting the 2-(5-Methyl-2-phenyl-4-oxazolyl)ethanol methanesulfonyl ester with 4-hydroxybenzothiophene to yield 4-[2-(benzo[b]thiophene-4-yloxy)-ethyl]-5-methyl-2-phenyl-oxazole,
- f) reacting the 4-[2-(benzo[b]thiophene-4-yloxy)-ethyl]-5-methyl-2-phenyl-oxazole with formaldehyde and HBr to yield 4-[2-(7-Bromomethyl-benzo[b]thiophen-4-yloxy)-ethyl]-5-methyl-2-phenyl-oxazole, and
- g) reacting the 4-[2-(7-Bromomethyl-benzo[b]thiophen-4-yloxy)-ethyl]-5-methyl-2-phenyl-oxazole with 2,4-thiazolidine to yield 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedione.
- 9. The process of claim 8, further comprising
 - h) converting the 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedione to Sodium 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedionate.
- 10. A process for the preparation of 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedione or Sodium 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedionate comprising the steps:
 - a) reacting methyl 3-oxovalerate with methyl orthoformate to yield methyl (E)-3-methoxy-2-pentenoate,
 - b) brominating the methyl (E)-3-methoxy-2-pentenoate to form methyl (E)-4-bromo-3-methoxy-pent-2-enoate,

- c) reacting the methyl (E)-4-bromo-3-methoxy-pent-2-enoate with benzamide to yield methyl 2-(5-methyl-2-phenyl-4-oxazolyl)acetate,
- d) reducing the methyl 2-(5-methyl-2-phenyl-4-oxazolyl)acetate to 2-(5-methyl-2-phenyl-4-oxazolyl)ethanol,
- e) reacting the 2-(5-methyl-2-phenyl-4-oxazolyl)ethanol with methanesulfonylchloride to yield 2-(5-methyl-2-phenyl-4-oxazolyl)ethanol methansulfonyl ester,
- f) reacting the 2-(5-Methyl-2-phenyl-4-oxazolyl)ethanol methanesulfonyl ester with 4-hydroxybenzothiophene to yield 4-[2-(benzo[b]thiophene-4-yloxy)-ethyl]-5-methyl-2-phenyl-oxazole,
- g) reacting the 4-[2-(benzo[b]thiophene-4-yloxy)-ethyl]-5-methyl-2-phenyl-oxazole with formaldehyde and HBr to yield 4-[2-(7-Bromomethyl-benzo[b]thiophen-4-yloxy)-ethyl]-5-methyl-2-phenyl-oxazole, and
- h) reacting the 4-[2-(7-Bromomethyl-benzo[b]thiophen-4-yloxy)-ethyl]-5-methyl-2-phenyl-oxazole with 2,4-thiazolidine to yield 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedione.
- 11. The process of claim 10, further comprising
 - i) converting the 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedione to Sodium 5-{4-[2-(5-Methyl-2-phenyl-oxazol-4-yl)-ethoxy]-benzo[b]thiophen-7-ylmethyl}2,4-thiazolidinedionate.

12. A compound of formula X

wherein

Y is Cl or Br,

R³ is lower alkyl, and

 R^4 is lower alkyl, lower-alkyl-carbonyl, lower alkoxy-carbonyl, aryl-carbonyl, $P(O)(OR^5)_2$ or $Si(R^6)_3$,

with the provisio that R⁴ may not be methyl if Y is Br or if R³ is methyl.
